Listing of Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

1. (original) A compound of formula (I)

$$\begin{array}{c} R^1 \\ O \\ R^2 \\ \hline \\ R^3 \\ X^1 = X^2 \end{array} \qquad \begin{array}{c} O \\ CH_2)_{\overline{n}} \\ \overline{C} - Y - R^5 \end{array} \qquad (I),$$

the *N*-oxides, the pharmaceutically acceptable acid addition salts and the stereochemically isomeric forms thereof, wherein

R¹ is hydrogen, C₁₋₄alkyl, halo, or polyhaloC₁₋₄alkyl;

R² is hydrogen, C₁₋₄alkyl, halo, or polyhaloC₁₋₄alkyl;

 R^3 is hydrogen or C_{1-4} alkyl;

R⁴ is hydrogen, C₁₋₄alkyl, or halo;

n is an integer 0, or 1;

 X^1 is carbon and X^2 is carbon; or X^1 is nitrogen and X^2 is carbon;

or X^1 is carbon and X^2 is nitrogen;

X³ is carbon or nitrogen;

Y represents O, or NR⁶ wherein R⁶ is hydrogen or C₁₋₄alkyl;

R⁵ represents a radical of formula

$$--(CH_2)_{\overline{m}} - \overset{R^8}{\underset{R}{\overset{O}{\subset}}} - \overset{C}{\underset{C}{\overset{}{\subset}}} - Z - R^9$$
 (a-1)

wherein

m is an integer 0, 1, or 2;

R⁹ is hydrogen, C₁₋₄alkyl, aryl¹, or C₁₋₄alkyl substituted with aryl¹;

- or when Y represents NR 6 the radicals R 5 and R 6 may be taken together with the nitrogen to which they are attached to form pyrrolidinyl substituted with C $_{1-4}$ alkyloxycarbonyl and optionally further substituted with hydroxy; or piperidinyl substituted with C $_{1-4}$ alkyloxycarbonyl;
- aryl is phenyl; phenyl substituted with one, two or three substituents each independently selected from C₁₋₄alkyl, C₁₋₄alkyloxy, halo, hydroxy, nitro, cyano, C₁₋₄alkyloxycarbonyl, trifluoromethyl, or trifluoromethoxy; or benzo[1,3]dioxolyl;
- aryl¹ is phenyl; phenyl substituted with one, two or three substituents each independently selected from C₁₋₄alkyl, C₁₋₄alkyloxy, halo, hydroxy, nitro, cyano, C₁₋₄alkyloxycarbonyl, trifluoromethyl, or trifluoromethoxy; and heteroaryl is imidazolyl, thiazolyl, indolyl, or pyridinyl.
- 2. (original) A compound as claimed in claim 1 wherein X¹, X² and X³ are carbon.
- 3. (original) A compound as claimed in claim 1 wherein R¹ is trifluoromethyl; R² is hydrogen; R³ is hydrogen; R⁴ is hydrogen; X¹, X² and X³ are carbon; n is the integer 1; Y represents NR⁶ wherein R⁶ is hydrogen or methyl; and R⁵ is a radical of formula (a-1) wherein m is the integer 0.

- 4. (original) A compound as claimed in claim 1 wherein R¹ is trifluoromethyl; R² is hydrogen; R³ is hydrogen; R⁴ is hydrogen; X¹, X² and X³ are carbon; n is the integer 1; Y represents NR⁶ wherein R⁶ is hydrogen or methyl; and R⁵ is a radical of formula (a-1) wherein m is the integer 1.
- 5. (original) A compound as claimed in claim 1 wherein R¹ is trifluoromethyl; R² is hydrogen; R³ is hydrogen; R⁴ is hydrogen; X¹, X² and X³ are carbon; n is the integer 1; Y represents NR⁶ wherein R⁶ is hydrogen or methyl; and R⁵ is a radical of formula (a-2) wherein m is the integer 1.
- 6. (previously presented) A compound as claimed in claim 1 wherein R¹ is trifluoromethyl; R² is hydrogen; R³ is hydrogen; R⁴ is hydrogen; X¹, X² and X³ are carbon; n is the integer 1; Y represents NR⁶ and R⁵ and R⁶ are taken together with the nitrogen to which they are attached to form pyrrolidinyl substituted with C₁₋₄alkyloxycarbonyl and optionally further substituted with hydroxy, or piperidinyl substituted with C₁₋₄alkyloxy-carbonyl.
- 7. (previously presented) A pharmaceutical composition comprising a pharmaceutically acceptable carrier and a therapeutically active amount of a compound as claimed in claim 1.
- 8. (previously presented) A process for preparing a pharmaceutical composition as claimed in claim 7 wherein a therapeutically active amount of a compound as claimed in claim 1 is intimately mixed with a pharmaceutically acceptable carrier.
- 9. (Withdrawn)

10 - 15. (Cancelled)